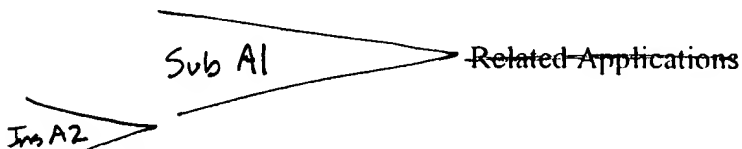# ENGRAFTABLE NEURAL PROGENITOR & STEM CELLS FOR BRAIN TUMOR THERAPY

FIELD OF THE INVENTION

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This invention is in the field of gene therapy, more particularly the field of using neuronal cells to treat brain tumors.

Sub A1   ~~Related Applications~~

10   In A2

This application is a continuation-in-part of pending U.S. Serial No. 09/133,873,
now U.S. Patent No. 5,958,767
filed on August 14, 1998, which is incorporated herein by reference.

Government Support

15

20   ## BACKGROUND

An effective gene therapy for the treatment of brain tumors has been an elusive goal for many years. Glioblastoma multiforma, which is virtually untreatable, and the less malignant anaplastic astrocytoma account for about one-quarter of the 5,000 intracranial

25   gliomas diagnosed yearly in the United States; 75 percent of gliomas in adults are of this category. Because of its profound and uniform morbidity, it contributes more to the cost of cancer on a per capita basis than does any other tumor. The patient, commonly stricken in the fifth decade of life, enters a cycle of repetitive hospitalizations and operations while experiencing the progressive complications associated with relatively

30   ineffective treatments of radiation and chemotherapy ("Harrison's Principles of Internal

Medicine," edited by Isselbacher, Braunwald, Wilson, Martin, Fauci and Kasper, 13th Edition, p.2262, McGraw-Hill, Inc. 1994).

One of the impediments to gene therapy of brain tumors such as gliomas, has been the degree to which they expand, migrate widely and infiltrate normal tissue. Most gene therapy strategies to date are viral vector-based, yet extensive distributions of sufficient amounts of viral vector-mediated genes to large regions and numbers of cells typically in need has often been disappointingly limited. Interestingly, one of the defining features of normal neural progenitors and stem cells is their migratory quality. Neural stem cells (NSCs) are immature, uncommitted cells that exist in the developing, and even adult, CNS and postulated to give rise to the array of more specialized cells of the CNS. They are operationally defined by their ability to self-renew and to differentiate into cells of most (if not all) neuronal and glial lineages in multiple anatomical & development contexts, and to populate developing and /or degenerating CNS regions.[1-5]

With the first recognition that neural cells with stem cell properties, reproduced in culture, could be reimplanted into mammalian brain where they could reintegrate appropriately and seamlessly in the neural architecture and stably express foreign genes[6-7], gene therapists began to speculate how such a phenomenon might be harnessed for therapeutic purposes. These, and the studies which they spawned (reviewed elsewhere[1-5,8]), provided hope that the use of neural progenitor/stem cells, by virtue of their inherent biology, might circumvent some of the present limitations of presently available gene transfer vehicles (e.g., non-neural cells, viral vectors, synthetic pumps), and provide the basis for a variety of novel therapeutic strategies.

Their use as graft material has been clearly illustrated by the prototypical neural progenitor clone, C17.2, a clone with which we have had extensive experience[6,9-16,17] and which was used in the studies presented here. C17.2 is a mouse cell line from postnatal day 0 cerebellum immortalized by infection with a retroviral construct containing the avian myc gene. This line has been transduced to constitutively express the *lacZ* and *neo*R genes. When transplanted into germinal zones throughout the brain, these cells have been shown to migrate, cease dividing, and participate in the normal development of multiple regions at multiple stages (fetus to adult) along the murine neuraxis,

differentiating appropriately into diverse neuronal and glial cell types as normal, non-tumorigenic cytoarchitectural constituents. They intermingle non-disruptively with endogenous neural progenitor/stem cells, responding to the same spatial and temporal cues in a similar manner. Crucial for therapeutic considerations, the structures to which

5   C17.2 cells contribute develop and maintain neuroanatomical normality. In their earliest therapeutic use, they served to deliver a missing gene product throughout the brains of mice with a lysosomal deficiency state and cross-corrected host cells by release and uptake of a lysosomal enzyme[9]. The feasibility of a neural progenitor/stem cell-based strategy for the delivery of therapeutic molecules directly to and throughout the CNS was

10   first affirmed by correcting the widespread neuropathology of a murine model of the genetic neurodegenerative lysosomal storage disease mucopolysaccaridosis type VII, caused by an inherited deletion of the β-glucuronidase (GUSB) gene, a condition that causes mental retardation and early death in humans. Exploiting their ability to engraft diffusely and become integral members of structures throughout the host CNS, GUSB-

15   secreting NSCs were introduced at birth into subventricular germinal zone, and provided correction of lysosomal storage in neurons and glia throughout mutant brains. In so doing, it established that neural transplantation of neural progenitor cells could provide a novel therapeutic modality.

What is needed is a way to treat tumors which are diffuse, infiltrating and/or

20   metastasizing. What is needed is a way to treat tumors locally to maximize the impact on the tumor and reduce the toxicity to the patient.

SUMMARY OF THE INVENTION

25   An isolated pluripotent neuronal cell having the capacity to differentiate into at least different types of nerve cells is disclosed. The pluripotent cell is further characterized by having a migratory capacity whereby the cell is capable of travelling from a first location where the neuronal cell is administered to a second location at which there is at least one tumor cell, having the ability to travel through and around a tumor,

30   whereby a plurality of the neuronal cells are capable of surrounding the tumor; and

having the capacity to track at least one infiltrating tumor cell, thereby treating infiltrating and metastasizing tumors.

The neuronal cell may be an isolated neural stem cell. The neuronal cell is optionally treated to secrete a cytotoxic substance. The neuronal cell alternatively is
5   transformed with factors that directly promote differentiation of neoplastic cells. Alternatively, the neuronal cell is transformed with viral vectors encoding therapeutic genes to be incorporated by tumor cells. In another embodiment, the neuronal cell can be transformed with viral vectors encoding suicide genes, differentiating agents, or receptors to trophins to be incorporated into tumor cells. The neuronal cells if
10   administered on the same side or a contralateral side of the brain from the tumor, are capable of reaching the tumor.

In another embodiment there is provided a method of converting a migrating neuronal cell to a migrating packaging/producer cell, said method includes the steps of a) providing a neuronal cell which constitutively produces a marker such as β-gal; b)
15   cotransfecting the neuronal cell with an amphotropic pPAM3 packaging plasmid and a puromycin selection plasmid pPGKpuro; c) selecting transfected cells in puromycin; d) selecting for cell surface expression of the amphotropic envelope glycoprotein coat; e) isolating cells by fluorescent activated cell sorting using monoclonal antibody 83A25; and f) screening the cells of step e for their packaging ability by assessing which colonies
20   packaged lacZ into infectious viral particles. Thus there is produced a migratory neuronal cell capable of being transfected with a gene of choice, so that viral particles expressing the gene of choice are produced and disseminated over a wide area of the central nervous system by a plurality of the transfected packaging cells.

The method of converting the migratory neuronal cell into a packaging cell line
25   wherein step f is performed by a virus focus assay for β-gal production. Alternatively the method can be performed with a prodrug activation enzyme as the gene of choice. Alternatively, the prodrug activation enzyme is E.coli cytosine deaminase (CD), HSV-TK or cytochrome p450. More preferably, the prodrug activation enzyme is E.coli cytosine deaminase (CD).

Also disclosed is a novel cell packaging line for the central nervous system. The cell line includes neuronal cells which constitutively produce a marker such as β-gal, have been cotransfected with an amphotropic pPAM3 packaging plasmid and a puromycin selection plasmid pPGKpuro; are selected in puromycin, for cell surface

5 expression of the amphotropic envelope glycoprotein coat and for fluorescence using monoclonal antibody 83A25, and for their packaging ability by assessing which colonies packaged lacZ into infectious viral particles. The resulting cells are capable of packaging and releasing particles or vectors which, in turn, may serve as vectors for gene transfer to central nervous system cells. The particles in the novel cell packaging line can be

10 replication-defective retroviral particles. The vectors in the novel cell packaging line can be replication-conditional herpes virus vectors.

BRIEF DESCRIPTION OF DRAWINGS

Figures 1A and 1B illustrate the migratory capacity of neural progenitor/stem

15 C17.2 cells *in vitro*. After 5 days of incubation there was a wide distribution of C17.2 cells (Fig. 1B), suggesting that they had migrated far from their initial seeding in the cylinder, compared to TR-10 cells (Fig. 1A), which remained localized to the area of initial seeding in the cylinders. These patterns were observed whether the cells were plated directly on top of the glioma cells (right-sided cylinder [arrows]) or simply in

20 juxtaposition to them (center cylinder [arrows]).

Figures 2A, 2B, 2C and 2D illustrate foreign gene-expressing neural progenitor/stem cells extensive migration throughout experimental tumor mass, and slightly beyond advancing tumor edge, appearing to "track" migrating tumor cells. (Fig. 2A) day 2 shown at 4X; arrowheads demarcate the approximate edges of tumor mass;

25 (Fig. 2B) high power at 10X where Xgal, blue-staining NSCs [arrows] are interspersed between tumor cells staining dark red. (Fig. 2C) View of tumor mass 10 days after intra-tumoral injection showing Xgal+blue, C17.2 NSCs have infiltrated the tumor but largely stop at the edge of the darkly red stained tumor tissue with some migration into surrounding tissue when the blue-staining NSC appears to be "following" an invading,

30 "escaping" cell [arrow] (10X). (Fig. 2D) CNS-1 tumor cells implanted into an adult

nude mouse frontal cortex, there is extensive migration and distribution of blue C17.2 cells throughout the infiltrating experimental tumor bed, up to and along the infiltrating tumor edge [arrows], and where many tumor cells are invading normal tissue, into surrounding tissue in virtual juxtaposition to aggressive tumor cells [arrows] (10X).

5   Figures 3A, 3B, 3C, 3D, 3E, 3F, 3G and 3H illustrate the neural progenitor/stem cells appearance to "track" migrating tumor cells away from main tumor mass; (Figs. 3A, 3B) parallel sections: low power C17.2 cells distributed throughout tumor and surrounding edge [Fig. 3A) Xgal and neutral red, Fig. 3B) double immunofluorescent labelling with texas red and FITC]; (Figs. 3C, 3D) low and high power of tumor edge and 10   migrating tumor cell in juxtaposition to C17.2 cell (Xgal and neutral red); (Figs. 3G, 3H) low and high power of single migrating tumor cells in juxtaposition to C17.2 cells (double immunoflourescent labelling with texas red and FITC).

Figures 4A, 4B, 4C, 4D, 4E, 4F and 4G illustrate neural progenitor/stem cells implanted at distant site from main tumor bed migrating throughout normal tissue target 15   CNS-1 tumor cells; (Figs. 4A, 4B) same hemisphere: 3x10^4 CNS-1 tumor cells implanted into right frontal lobe. On day 6, 4x10^4 C17-2 cells injected into right frontoparietal lobe (4mm caudal tumor injection). Animals sacrificed on day 12 (shown) and day 21, C17-2 cells seen in tumor bed (Xgal and neutral red). (Figs. 4C, 4D, 4E) Contralateral hemisphere: 3x10^4 CNS-1 tumor cells implanted into left frontal lobe and 5x10^4 CNS-1 20   tumor cells implanted into left frontoparietal lobe. On day 6, 8x10^4 C17-2 cells were injected into right front lobe. Animals were sacrificed on day 12 and 21 (shown); c) 4x C17.2 cells (red) seen migrating towards tumor (green) from opposite side of the brain, d) 10x C17.2 cells (red) seen actively migrating across central commisure (double immunofluorescence), e) 20x C17-2 cells (blue) seen entering tumor (black arrows) 25   (Xgal/ neutral red). (Figs. 4F, 4G) Intraventricular 5x10^4 CNS-1 tumor cells were implanted into right frontal lobe. On day 6, 8x10^4 C17.2 cells were injected into right or left (shown) lateral ventricle.

DETAILED DESCRIPTION OF INVENTION

The experiments presented herein demonstrate that NSCs (prototypical clone C17.2) when implanted into an experimental glioma, will distribute throughout the tumor and migrate along with aggressively advancing tumor cells, while continuing to express

5   their reporter gene lacZ. (One of the glioma lines used, astrocytoma cell line CNS-1, demonstrates single cell infiltration and invasive characteristics similar to those of human glioblastomas[18]). Furthermore, the neural progenitor/stem cells seem to migrate slightly beyond and surround the invading tumor border. In additional experiments, where neural progenitors were implanted at a distant site from the tumor bed, in the same hemisphere,

10   opposite hemisphere, or lateral ventricle, they migrated through normal tissue moving specifically toward CNS-1 tumor cells. They were found to accumulate in or near the tumor bed as well as near or in direct juxtaposition to the individual infiltrating tumor cells.

Not wishing to be bound by any particular theory, the inventors propose that this

15   neural progenitor/stem cell system migrate towards a trophic gradient of growth factors produced by the tumor cells. Thus, NSCs may provide a unique platform for the dissemination of therapeutic genes to the proximity of or into tumors that previously were inaccessible. These observations further suggest a number of other new gene therapy approaches. These may include the dissemination of cytotoxic gene products, but could

20   also include factors that directly promote differentiation of neoplastic cells as well as the more efficacious delivery of viral vectors encoding therapeutic genes to be incorporated by tumor cells (e.g. suicide genes, differentiating agents, receptors to trophins). Because NSCs can be engineered to package and release replication-defective retroviral particles or replication-conditional herpes virus vectors which, in turn, may serve as vectors for the

25   transfer of genes to CNS cells, neural progenitor/stem cells should serve to magnify the efficacy of viral-mediated gene delivery to large regions in the brain.

One effective mode of therapy for experimental brain tumors has been prodrug activation. Initially, prodrug activation enzymes were limited to antibodies directed against tumor enriched antigens. New strategies incorporate genes for these enzymes into

30   viral vectors. Among the prodrug activating systems shown to be effective for gliomas

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E.coli cytosine deaminase (CD), HSV-TK and cytochrome p450 have been demonstrated to have a drug mediated bystander effect. Of these CD gives the best reported "bystander" effect. CD converts the nontoxic prodrug 5-fluorocytosine (5-FC) to 5-fluorouridine (5-FU) metabolites. 5-FU is a chemotherapeutic agent which has selective

5   toxicity for actively dividing cells, thus primarily targeting tumor cells. In addition, 5-FU and its toxic metabolites can readily pass into adjacent and surrounding cells by non-facilitated diffusion. Brain tumors may require only a small number of cells expressing CD (about 2% evenly distributed) to generate significant anti-tumor effects when treated with systemic, non-toxic levels of 5-FC. Our results support the hypothesis that

10   transduced NSCs would disperse CD expression efficiently throughout the tumor and even "track" single migrating, "escaping" tumor cells.

Another approach to brain tumor gene therapy has been selective gene transfer to tumor cells in combination with pharmacotherapy, e.g., the HSV-TK gene, when transduced via retrovirus into a dividing population of brain tumor cells, confers a lethal

15   sensitivity to the drug ganciclovir. Recent modifications of retroviral constructs to increase efficiency of infection and cell-specific targeting hold promise for enhancing the potency of this strategy. Again, through the "bystander effect", tumor destruction is effective even when only a fraction of the cells express HSV-TK; adjacent tumor cells not expressing HSV-TK also appear to be eliminated. Attempts to improve efficiency of

20   tumor destruction have focused on increasing the number of cells expressing the HSV-TK gene. The use of NSCs as packaging cells (which might then be self-eliminated) may prove to be an effective extended delivery system of the lethal gene to neighboring mitotic tumor cells, especially individual, infiltrating tumor cells.

In conclusion, genetically modified neural progenitor/stem cells have the potential

25   to supply a range of tumor selective agents throughout mature and developing brains. The experiments presented here demonstrate the ability of NSCs: (1) to migrate/distribute quickly and effectively throughout the main tumor bed when implanted directly into the experimental gliomas; (2) to migrate slightly beyond and "surround" (as if to contain) the invading tumor border; (3) to seemingly "track" individual, infiltrating tumor cells into

30   surrounding tissue; (4) to migrate through normal tissue from distant sites to target CNS-

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1 tumors; and (5) to show stable expression of a foreign gene, in this case *lacZ*, throughout the tumor bed and in juxtaposition to tumor cells. These results lay the groundwork for future therapeutic brain tumor studies, providing critical support for the use of neural progenitor/stem cells as an effective delivery vehicle for tumor directed,

5   vector-mediated enzyme/prodrug gene therapy.

Other cells

The HCN-1 cell line is derived from parental cell lines from the cortical tissue of a patients with unilateral megalencephaly growth (Ronnett G.V. etal. Science 248:603-5, 1990). HCN-1A cells have been induced to differentiate to a neuronal-like morphology

10   and stain positively for neurofilament, neuron-specific enolase and p75NGFR, but not for myelin basic protein, S-100 or glial fibrillary acidic protein (GFAP). Because these cells also stain positively for γ-amino butyric acid and glutamate, they appear to become neuro-transmitting bodies. Earlier Poltorak M et al. (Cell Transplant 1(1):3-15, 1992) observed that HCN-1 cells survived in the brain parenchyma and proposed that these cells

15   may be suitable for intracerebral transplantation in humans.

Ronnet GV et al. (Neuroscience 63(4):1081-99, 1994) reported that HCN-1 cells grew processes resembling neurons when exposed to nerve growth factor, dibutyryl cyclic AMP and isobutylmethylxanthine.

The nerve cells also can be administered with macrophages which have been

20   activated by exposure to peripheral nerve cells. Such activated macrophages have been shown to clean up the site of CNS trauma, for example a severed optic nerve, after which new nerve extensions started to grow across the lesion. Implanting macrophages exposed to CNS tissue (which secretes a chemical to inhibit macrophages) or nothing at all resulted in little or no regeneration (Lazarov-Spiegler et al. FASEB J. 10:1, 1996).

25   Fetal pig cells have been implanted into patients with neurodegenerative diseases, such as Parkinson's disease and Huntington's chorea, and intractable seizures, in whom surgical removal of the excited area would otherwise have been performed. Such cells, if properly screened for retroviruses, could also be used in the inventive method.

Neural crest cells are isolated and cultured according to Stemple and Anderson

30   (U.S. Patent No. 5,654,183), which is incorporated herein by reference, with the

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modification that basic fibroblast growth factor (bFGF) is added to the medium at

concentrations ranging from 5 to 100 ng/ml in 5 ng/ml increments. Neural crest cells so

cultured are found to be stimulated by the presence of FGF in increasing concentrations

about 1 or 5 ng/ml. Such cells differentiate into peripheral nerve cells, which can be used

5   in the instant invention.

Other cytokines, growth factors and drugs

Certain cytokines, growth factors and drugs are optionally used in the trnasplant

area or may be administered concomitantly with the transplant.

Known cytokines include interleukins (IL) IL-1, IL-2, IL-3, IL-4, IL-5, IL-6, IL-7,

10   IL-8, IL-10, and IL-11; tissue necrosis factors (TNF), TNFα, also lymphotoxin (LT) and

TNFβ; interferons (IFN) IFNα, IFNβ and IFNγ; and tissue growth factor (TGF). The

colony-stimulating factors (CSFs) are specific glycoproteins that are thought to be

involved in the production, differentiation and function of stem cells.

Nerve growth factor (NGF) has been shown to increase the rate of recovery in

15   spatial alternation tasks after entorhinal lesions, possibly by acting on cholinergic

pathways (Stein and Will, Brain Res. 261:127-31, 1983).

<div align="center">EXAMPLES</div>

20   Experimental Methods

Cells:

C17-2 and TR-10 cells were maintained in Dulbecco's modified Eagle's medium

(DMEM; Mediatech, Washington, DC) supplemented with 10% fetal calf serum (FCS;

Sigma, St. Louise, MO), 5% horse serum (HS; Gibco), 1% Glutamine (2mM; Gibco), 1%

25   penicillin/streptomycin (Sigma). CNS-1 cells were stably transduced with the PGK-GFP-

IRES-NeoR retroviral vector construct to express green fluorescent protein (GFP) as

previously described (ref. Aboody-Guterman et. al, 1997), and maintained in RPMI-1640

(Bio Whittaker) supplemented with 10% FCS and 1% penicillin/streptomycin (Sigma).

Cell structure studies were performed in 100mm petri dishes under standard conditions:

30   humidified, 37°C, 5% CO_2 incubator. *In vitro* studies; CNS-1 glioma cells were plated to

<div align="center">10</div>

approximately 60-70% confluency around a 5mm cylinder (i.e. free of CNS-1 cells) into which 40,000 C17.2 or TR-10 cells plated overnight. At the same time, 40,000 C17.2 or TR-10 cells were placed into a 5mm cylinder placed directly on top of adhered CNS-1 cells. The next day, cylinders were removed and plates rinsed well with PBS to remove

5   any floating cells, media was replaced, and plates incubated for 5 days. Plates were subsequently stained for β-galactosidase overnight after .5% glutaraldehyde fixation. (Note: both C17.2 and TR-10 cells are >90% blue with X-gal staining). *In vivo* studies; 48 hours prior to transplant, C17.2 and TR-10 cells were incubated with BUdR (Sigma) at a concentration of 10μM. Plated cells were rinsed with PBS, trypsinized, resuspended

10   in media and counted on the Coulter counter. Desired number of cells were spun down at 4°C in the centrifuge for 4 minutes and 1100 rpm to obtain a pellet. Media was removed; cells were rinsed by resuspending in PBS and respun. PBS was removed and the appropriate amount of PBS added to resuspend cells at final desired concentration. Cells were kept on ice, and gently triturated prior to each animal injection. Cells not labelled

15   with BUdR were prepared for injection in similar manner.

Animals:

Animal studies were performed in accordance with guidelines issued by the Massachusetts General Hospital Subcommittee on Animal Care. Animals used: adult CD-Fisher rats (Charles River) and 8-10 week old adult, approximately 20 gram female

20   nude mice (random bred Swiss white obtained from Cox 7, MGH-East).

Surgery and Sacrifice:

Animals were anesthetized by an i.p. injection of .15 ml of 20% ketamine HCL (KETALAR 100 mg/ml; Parke-Davis, Morris Plains, NJ), 20% xylazine (ROMPUN 20 mg/ml; Miles Inc., Shawnee Mission, KS), 60% sodium chloride (0.9%; Abbott

25   Laboratories, North Chicago, IL) and immobilized in stereotactic apparatus (Kopf, Tujunga, CA). Intracerebral injections were stereotactically performed by making a linear scalpel skin incision on top of the skull. A burr hole was drilled into the skull with a high speed drill 2mm lateral to the bregma on the coronal suture. After incising the dura with a sterile needle and obtaining hemostasis, desired number on tumor cells

30   suspended in 1 μl of 1X Dulbecco's phosphate-buffered salt solution (PBS pH 7.4;

Mediatech, Herndon, VA) were injected with a 26 guage 5 µl Hamilton syringe to specified location (see protocols below) over a 3 to 5 minute period. After retracting the needle over a 2-4 minute period, bone-wax (Ethicon, Somerville, NJ) was used to occlude the burr hole, betadine applied to surgical area, and the skin sutured closed. Animals

5   receiving a second injection at a later date were anesthetized, immobilized in stereotactic apparatus, and cells injected as per specific protocol (see below). Animals were sacrificed on stated days with an overdose of anesthesia and subsequent intracardiac perfusion with PBS followed by 4% paraformaldehyde+2mM $MgCl_2$ (pH 7.4). Brains were removed and post-fixed overnight at 4 °C and then transferred to 30% sucrose in

10   PBS + 2mM $MgCl_2$ (pH 7.4) for 3-7 days to cryoprotect. Brains were stored at -80 °C and then 10-15 micron coronal serial sections were cut to cryostat (Leica CM 3000).

Sub B2 ————> ~~BUdR labelling of engrafted C17-2 cells~~:

Selected animals received 3 intraperitoneal injections of 1 ml/100g body weight 20uM BUdR stock solution (Sigma) over 24 hours prior to sacrifice (.2 ml/injection per

15   20g mouse).

Histopathological and Immunohistochemical Studies:

Tissue sections were stained with (1) X-gal and counterstained with neutral red (2) hematoxylin and eosin (3), double immunofluorescent labelling was performed with texas red anti-beta-galactosidase and FITC anti-GFP. Slides were examined with light

20   microscopy, fluorescent microscopy. CNS-1 tumor cells were also examined without staining under confocal fluorescent microscopy.

Example 1. Migratory Capacity of NSCs in Culture

To determine properties of the NSCs in association with glioma cells, studies

25   were initially performed in culture comparing the relative migratory capacity of NSCs (clone C17.2) to fibroblasts (the lacZ-expressing TR-10 fibroblast cell line) when co-cultured with glioma cells. C17.2 and TR-10 cells were maintained in Dulbecco's modified Eagle's medium (DMEM; Mediatech, Washington, DC) supplemented with 10% fetal calf serum (FCS; Sigma, St. Louise, MO), 5% horse serum (HS;Gibco), 1%

30   Glutamine (2mM; Gibco), 1% penicillin/streptomycin (Sigma). CNS-1 cells were stably

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transduced with the PGK-GFP-IRES-NeoR retroviral vector construct to express green

fluorescent protein (GFP) as previously described (ref. Aboody-Guterman et. al, 1997),

and maintained in RPMI-1640 (Bio Whittaker) supplemented with 10% FCS and 1%

penicillin/streptomycin (Sigma). Cell structure studies were performed in 100mm petri

5   dishes under standard conditions: humidified, 37 ° C, 5% CO2 incubator. CNS-1 glioma

cells were plated to approx. 60-70% confluency around a 5mm cylinder (i.e. free of CNS-1

cells) into which 40,000 C17.2 or TR-10 cells plated overnight. At the same time, 40,000

C17.2 or TR-10 cells were placed into a 5mm cylinder placed directly on top of adhered

CNS-1 cells. The next day, cylinders were removed and plates rinsed well with

10   PBS to remove any floating cells, media was replaced, and plates incubated for 5 days.

Plates were subsequently stained for β-galactosidase overnight after .5% glutaraldehyde

fixation. (Note: both C17.2 and TR-l0 cells are >90% blue with X-gal staining).

There was a wide distribution of C17.2 cells (Fig. 1B), suggesting that they had

migrated far from their initial sites in the cylinder, compared to the T-10 cells (Fig. lA),

15   which remained localized to the area of initial seeding in the cylinders. These patterns were

observed whether the cells were plated directly on top of the glioma cells (right-sided

cylinder [arrows]) or simply in juxtaposition to them (center cylinder [arrows]).

Example 2. Transgene-Expressing NSCs Migrate Throughout and Beyond Invading

20   **Tumor Mass *in vivo***

To determine the behavior of clone C17.2 NSCs introduced into brain tumors,

experimental animals (syngeneic adult rats) first received an implant of 4×10^4 D74 rat

glioma cells in 1 μl injected into the right frontal lobe. Four days later, 1×10^5 C17.2 NSCs

in 1.5 μl PBS were injected at same coordinates directly into the D74 tumor bed.

25   Animals were then sacrificed at days 2, 6, and 10 post-intratumoral injection and cryostat

sections of the brains were processed with Xgal histochemistry for β-galactosidase (βgal)

activity to detect donor-derived cells and counterstained with neutral red to detect tumor

cells.

Donor C17.2 NSCs were found extensively dispersed throughout the tumor

30   within a few days, spanning an 8mm width of tumor as rapidly as 2 days after injection (Figs.

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transduced with the PGK-GFP-IRES-NeoR retroviral vector construct to express green fluorescent protein (GFP) as previously described (ref. Aboody-Guterman et. al, 1997), and maintained in RPMI-1640 (Bio Whittaker) supplemented with 10% FCS and 1% penicillin/streptomycin (Sigma). Cell structure studies were performed in 100mm petri

5 dishes under standard conditions: humidified, 37 ° C, 5% CO_2 incubator. CNS-1 glioma cells were plated to approx. 60-70% confluency around a 5mm cylinder (i.e. free of CNS-1 cells) into which 40,000 C17.2 or TR-10 cells plated overnight. At the same time, 40,000 C17.2 or TR-10 cells were placed into a 5mm cylinder placed directly on top of adhered CNS-1 cells. The next day, cylinders were removed and plates rinsed well with

10 PBS to remove any floating cells, media was replaced, and plates incubated for 5 days. Plates were subsequently stained for β-galactosidase overnight after .5% glutaraldehyde fixation. (Note: both C17.2 and TR-10 cells are >90% blue with X-gal staining).

There was a wide distribution of C17.2 cells (Fig. 1B), suggesting that they had migrated far from their initial sites in the cylinder, compared to the TR-10 cells (Fig. 1A),

15 which remained localized to the area of initial seeding in the cylinders. These patterns were observed whether the cells were plated directly on top of the glioma cells (right-sided cylinder [arrows]) or simply in juxtaposition to them (center cylinder [arrows]).

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25 Animals were then sacrificed at days 2, 6, and 10 post-intratumoral injection and cryostat sections of the brains were processed with Xgal histochemistry for β-galactosidase (βgal) activity to detect donor-derived cells and counterstained with neutral red to detect tumor cells.

Donor C17.2 NSCs were found extensively dispersed throughout the tumor within
30 a few days. spanning an ~8mm width of tumor as rapidly as 2 days after injection (Figs.

2A, 2B). This is a much more extensive and rapid dispersion compared to previous reports of 3T3 fibroblasts grafted into an experimental brain tumor[33]. By day 10, C17.2 cells were seen throughout a majority of the tumor, clearly along the infiltrating tumor edge and slightly beyond it, drawn somewhat by the degenerative environment, seeming

5 to "track" migrating tumor cells (Figs. 2C, 2D). C17.2 cells themselves did not become tumorigenic.

(Fig. 2A) Day 2 shown at 4X; arrowheads demarcate the approximate edges of tumor mass; even at lower power, the tumor can be seen to be intermixed with blue NSCs [arrows]. This is appreciated more dramatically at high power in (Fig. 2B) at 10X where

10 Xgal+, blue-staining NSCs [arrow] are interspersed between tumor cells staining dark red. (Fig. 2C) This view of the tumor mass, 10 days after intra-tumoral injection nicely shows that Xgal+blue, C17.2 NSCs have infiltrated the tumor but largely stop at the edge of the darkly red stained tumor tissue (border indicated by arrowheads) with some migration into surrounding tissue when blue-staining NSC appears to be "following" and

15 invading, "escaping" tumor cell [arrow] (10X). This phenomenon becomes even more dramatic when examining the behavior of C17.2 NSCs in an even more virulent, invasive and aggressive tumor than D74, the experimental CNS-1 astrocytoma in the brain of a nude mouse (Fig. 2D). CNS-1 tumor cells were implanted into an adult nude mouse frontal cortex (day 0). On day 6, 4×10^4 C17.2 cells were implanted directly into the

20 tumor bed. The animal pictured in (Fig. 2D) was sacrificed on day 12 post-tumor implantation, 6 days post-intra-tumoral injection. The cryostat section pictured was processed with Xgal histochemistry for β-galactosidase activity to detect blue C17.2 NSCs and counterstained with neutral red to show dark red tumor cells. There is extensive migration and distribution of blue C17.2 cells throughout the infiltrating

25 experimental tumor bed, up to and along the infiltrating tumor edge [white arrows], and, where many tumor cells are invading normal tissue, into surrounding tissue in virtual juxtaposition to aggressive tumor cells [arrows] (10X).

Example 3. NSCs "Track" Infiltrating Tumor Cells

CNS-1 tumor cells were labelled by retroviral transduction with green fluorescent protein (GFP), prior to implantation, to better distinguish single cells away from the main tumor bed[17]. GFP-expressing CNS-1 glioma cells (3 x 10⁴) in 1 µl PBS injected into right frontal lobe at stereotaxic coordinates 2mm lateral to bregma, on coronal suture, 3mm

5   depth from dura. 4 x 10⁴ C17.2 or TR-10 cells in 1 µl PBS injected at same coordinates directly into tumor bed on day 6. 3-4 C17.2 animals (2 BUdR labelled, 1 BUdR pulsed) and 1-2 TR-10 control animals (1 BUdR labelled). Animals were sacrificed on days 9,12, 16 and 21 post-tumor implantation. Cryostat sectioned, fixed brain tissue was stained either with β-galactosidase (C17.2 cells blue) and neutral red (tumor cells dark red) or

10   double immunofluorescence with Texas red anti-β-galactosidase (C17.2 cells red) and FITC anti-GFP (tumor cells green).

(Figs. 3A, 3B) parallel sections: low power of C17.2 cells distributed throughout tumor and surrounding edge [Fig.3A) Xgal and neutral red, Fig.3B) double immunofluorescent labelling with Texas red and FITC]

15   (Figs. 3C, 3D) low and high power of single migrating tumor cell in juxtaposition to C17.2 cell (Xgal and neutral red)

(Figs. 3E, 3F) low and high power of single migrating tumor cell in juxtaposition to C17.2 cell (Xgal and neutral red)

(Figs. 3G, 3H) low and high power of single migrating tumor cells in juxtaposition to

20   C17.2 cells (double immunofluorescent labelling with Texas red and FITC).

Example 4. NSCs Implanted at Distant Site Migrate Toward Tumor

To examine the capacity of NSCs to migrate through normal tissue and specifically target tumor cells, donor NSCs were injected into uninvolved sites distant

25   from the main tumor bed in three separate paradigms, into the same hemisphere, into the opposite hemisphere, or into the lateral ventricles.

Same hemisphere: CNS-1 glioma cells (3 x 10⁴) in 1 µl PBS was injected into the right frontal lobe at stereotaxic coordinates 2mm lateral to bregma, on coronal suture, 3mm depth from dura. 4 x 10⁴ C17.2 or TR-10 cells in 1 µl PBS injected into right frontal

30   parietal lobe at stereotaxic coordinates 3mm lateral and 4mm caudal to bregma, 3mm

depth from dura on day 6. Two animals were sacrificed at days 12 and 21. At all time points, NSCs were found distributed within the main tumor bed as well as in juxtaposition to migrating tumor cells in surrounding tissue (Figs. 4A, 4B).

Opposite hemisphere: 3 x 10^4 CNS-1 tumor cells in 1 μl PBS injected into left

5   frontal lobe at stereotaxic coordinates 2mm lateral to bregma, on coronal suture, 3mm depths from dura, 5 x 10^4 CNS-1 tumor cells in 1 μl PBS injected into left frontoparietal lobe 3mm lateral and 4mm caudal to bregma, 3mm depth from dura, 8 x 10^4 C17.2 cells in 2 μl PBS injected into right frontal lobe 2mm lateral and 2mm caudal to bregma, 3mm depth from dura on day 6. Two animals sacrificed on day 12 and 21. (control - no tumor

10   Coordinates: 2mm R of bregma, 2mm caudal, 3mm deep). NSCs were seen actively migrating across the central commissure towards the tumor on the opposite side of the brain, and then entering the tumor (Figs. 4C, 4D, 4E).

Implantation away from CNS-1 tumor bed (intraventricular):

In this final paradigm 5 x 10^4 CNS-1 tumor cells in 1 μl PBS was injected into the

15   right frontal lobe 2mm lateral to bregma, on coronal suture, 3mm depth from dura. 8 x 10^4 C17.2 cells in 2 μl PBS injected into left or right ventricle 1mm lateral and 3mm caudal to bregma, 2mm depth from dura on day 6. Two animals sacrificed on days 12 and 21. NSCs again were seen within the main tumor bed, as well as in juxtaposition to migrating tumor cells (Figs. 4F, 4G).

20   In each case, donor NSCs were found to migrate through normal tissue and "target" the tumor.

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